	Diagnosis							OI	JADAS	S 2 Iten	ns*						G	RADE (outcom	e level)		
Index test	Reference	uries Study	Likelihood	1	2	3	4	5	6	7	8	9	10	11	Study	Risk of	Indirectness	Inconsistency	Imprecise	Publication	Downgrade
	standard		ratio												design	bias		,	evidence ✓	bias	
Aktiv slump	MRI		LR+	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	√	✓	✓	×	?	V	?	↓
			LR-																√		\
Dain during CLD	MRI		LR+	√	✓	√	✓	✓	✓	✓	√	✓	√	✓	/	✓	×	?	✓	2	↓
Pain during SLR	WIKI		LR-	V	•	•	•	•	•	•	V	•	V	•	•	•		f	✓	- · ·	\
			LR+																✓		\
Pai during 90deg R KF	MRI		LR-	✓	✓	✓	√	√	✓	✓	✓	✓	\checkmark	✓	✓	✓	×	?	✓	?	\
																			✓		
Pai during 30deg R KF	MRI	Wangensteen et al. (1)	LR+	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	✓	×	?		?	\
		5 t and (1)	LR-																✓		\
Pain during active KF	MRI		LR+	√	✓	√	✓	✓	✓	✓	√	✓	✓	✓	✓	✓	✓	?	✓	· ?	\leftrightarrow
r air during active iti	IVII (I		LR-	•	·	·	•	·	ľ		·		·	•	·	•	•	•	✓	•	\leftrightarrow
			LR+																✓		\
Pain during active KE	MRI		LR-	√	✓	✓	✓	✓	✓	✓	√	√	✓	✓	✓	√	×	?	✓	?	\
			LR+																✓		\
Pain during trunk F	MRI		LR-	✓	✓	\checkmark	✓	✓	✓	✓	✓	✓	\checkmark	✓	✓	✓	×	?	√	?	\
Taking off shoe	US		LR+: N/A	×	✓	?	×	✓	✓	×	×	✓	?	?	✓	xx	×	?	N/A	?	N/A
Designed range of			LR- LR+: N/A																? N/A		↓↓↓ N/A
Resisted range of motion test	US	Zeren et al. (2)	LR-	×	√	?	×	√	✓	×	×	✓	;	?	√	xx	×	?	√ ×	?	→ → →
Passive range of motion test	US	(←)	LR+: N/A LR-	×	✓	?	×	✓	√	×	×	✓	?	?	✓	xx	×	?	N/A ✓	?	N/A ↓↓↓
Active range of motion	US		LR+: N/A	×	✓	?	×	√	✓	×	×	✓	2	?	/	xx	×	?	N/A	?	N/A
test	05		LR-	^	•	ſ	^	Y	V	^	^	V	?		•	~ ~	^	ſ	✓		$\downarrow\downarrow\downarrow\downarrow$
Composit	MRI	Schneider- Kolsky et al.	LR+	√	✓	√	✓	✓	✓	✓	√	✓	?	✓	✓	✓	×	?	✓	. ?	\
σοπροσιι	INIUI	(3)	LR-: N/A	•		•					•		ŗ						N/A	ŗ.	N/A

Abbreviations: MRI (magnetic resonance imaging); US (ultrasound); LR+ (Positive likelihood ratio); LR- (negative likelihood ratio); N/A (not applicable)

*Item 1: Was a consecutive or random sample of patients enrolled? Item 2: Was a case-control design avoided? Item 3: Did the study avoid inappropriate exclusions? Item 4: Were the index test results interpreted without knowledge of the results of the reference standard? Item 5: If a threshold was used, was it pre-specified? Item 6: Is the reference standard likely to correctly classify the target condition? Item 7: Were the reference standard results interpreted without knowledge of the results of the index test? Item 8: Was there an appropriate interval between index test(s) and reference standard? Item 9: Did all patients receive a reference standard? Item 10: Did patients receive the same reference standard? Item 11: Were all patients included in the analysis?

Quadas 2 risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: \times = item cause for possible downgrade once; \times = item cause for possible downgrade twice; \checkmark = item fulfilled, no downgrading; ? = item unclear or not available, no upgrading or downgrading.

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Table 2. Risk of bias assessment and GRADE of clinical tests for diagnosing adductor injuries.

	Diagnosi	is						_	NIAD/	AS Items							G	GRADE (outcon	ne level)		
	Adductor inj	uries						C	OADA	to items	5							ATTABL (OUTCOIL	ie ievei)		
Index test	Reference standard	Study	Likelihood ratio	1	2	3	4	5	6	7	8	9	10	11	Study design	Risk of bias	Indirectness	Inconsistency	Imprecise evidence	Publication bias	Downgrade**
Delegation	MRI		LR+	→	✓	✓	✓	✓	✓	√	✓	✓	ç	✓	✓	✓	×	?	✓	?	V
Palpation	IVINI		LR -			•	•	•	•	•	•	•	ŗ	•	•	•	^	· ·	x	ŗ	\
0 00	MDI		LR+	✓	✓	√	√	✓	√	√	√	√	?	✓	✓	✓		?	x	2	↓↓
Squeeze 0°	MRI		LR -		•	V	•	•	•	V	•	•	· ·	v	•	•	×	f	✓	?	V
0 450	MDI		LR+	√	✓	√	✓	✓	√	√		✓	2	✓				?	√	2	
Squeeze 45°	MRI	Serner et	LR -		•	V	•	•	•	V	√	•	?	V	√	✓	×	f f	•	?	V
Isometric adduction		al. (4)	LR+	✓	✓	√	✓	✓	√	√		✓	?	✓		✓		?	x		↓↓
(outer range)	MRI		LR -		•	V	•	•	•	V	√	•	?	V	√	•	×	f f	✓	· ?	V
			LR+	√			✓	✓	√			✓	?	✓				?	x	2	↓↓
Adductor stretching	MRI		LR -		√	√	V	V	•	√	√	V	?	V	√	✓	×	f f	✓	· ?	V
Flexion Abduction			LR+						√												
External Rotation (FABER)	MRI		LR -	- ✓	√	√	✓	✓	•	✓	√	✓	?	✓	√	✓	×	?	✓	?	V

Abbreviations: MRI (magnetic resonance imaging); LR+ (Positive likelihood ratio); LR- (negative likelihood ratio).

*Item 1: Was a consecutive or random sample of patients enrolled? Item 2: Was a case-control design avoided? Item 3: Did the study avoid inappropriate exclusions? Item 4: Were the index test results interpreted without knowledge of the results of the reference standard? Item 5: If a threshold was used, was it pre-specified? Item 6: Is the reference standard likely to correctly classify the target condition? Item 7: Were the reference standard results interpreted without knowledge of the results of the index test? Item 8: Was there an appropriate interval between index test(s) and reference standard? Item 9: Did all patients receive a reference standard? Item 10: Did patients receive the same reference standard? Item 11: Were all patients included in the analysis?

Quadas 2 risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: x = item cause for possible downgrade once; xx = item cause for possible downgrade twice; x = item fulfilled, no downgrading; ? = item unclear or not available, no upgrading or downgrading.

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Table 3. Risk of bias assessment and GRADE of clinical tests for diagnosing rectus femoris injuries.
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	Diagnosis	S						0		S Items							0	DADE (outcom	o lovol)		
Rec	tus femoris	injuries						G	OADA	S Items	•						G	RADE (outcom	e ievei)		
Index test	Reference standard	Study	Likelihood ratio	1	2	3	4	5	6	7	8	9	10	11	Study design	Risk of bias	Indirectness	Inconsistency	Imprecise evidence	Publication bias	Downgrade **
Palpation	MRI		LR+	✓	✓	√	/	✓	✓	√	/	1	2	✓	✓	✓	×	?	×	?	\
Ιαιρατίστι	IVII ti		LR -	·	·	·	,	•	,	·	•	,	:	·		•	_	:	?	:	\
Isometric hip flexion	MRI		LR+	\	✓	√	/	✓	✓	√	✓	✓	2	✓	✓	✓	×	?	✓	?	\
0°	IVINI		LR -	•	•	·	·	•	•	•	•	,	:	,		•	^	:	×	:	$\downarrow \downarrow$
Isometric hip flexion	MRI		LR+	\	<	√	✓	✓	✓	√	✓	✓	?	✓	✓	✓	×	?	✓	· ?	\
90°	IVINI		LR -		•	•	•	•	•	•	•	•		•	•	•	^	f	×	ŗ.	44
Isometric hip flexion (modified Thomas	MRI	Serner et	LR+	\	<	√		✓	✓	√	✓	✓	?	✓	✓	✓	×	?	✓	?	\
Test)	IVINI	al. (4)	LR -		•	•	•	•		•		,		,		•	^	f	×	ŗ.	44
Isometric knee extension (modified	MRI		LR+	✓	<	√	✓	✓	✓	√	✓	\	?	✓	✓	✓	×	?	✓	· ?	\
Thomas Test)	IVINI		LR -		•	•	•	•		•		,		,		•	^	f	?	ŗ.	\
Hip extension (stretching; modified	MRI		LR+	✓	✓	√	-/	✓	✓	√	✓	✓	?	√	✓	✓	×	?	✓	?	V
Thomas Test)	IVINI		LR -		•	•	•	•		•	•	,	:	,		•	^	f	×	ŗ	1
Knee flexion (stretching; modified	MRI		LR+	✓	✓	√	1	✓	✓	√	√	✓	· ·	√	✓	✓	×	?	×	?	\
Thomas Test)	IVIDI		LR -			,				,						_	^	· ·	×	· ·	\

Abbreviations: MRI (magnetic resonance imaging); LR+ (Positive likelihood ratio); LR- (negative likelihood ratio).

*Item 1: Was a consecutive or random sample of patients enrolled? Item 2: Was a case-control design avoided? Item 3: Did the study avoid inappropriate exclusions? Item 4: Were the index test results interpreted without knowledge of the results of the reference standard? Item 5: If a threshold was used, was it pre-specified? Item 6: Is the reference standard likely to correctly classify the target condition? Item 7: Were the reference standard results interpreted without knowledge of the results of the index test? Item 8: Was there an appropriate interval between index test(s) and reference standard? Item 9: Did all patients receive a reference standard? Item 10: Did patients receive the same reference standard? Item 11: Were all patients included in the analysis?

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Treatment Hamstring			Risk o	of Bias	asses	ssmer	it Item*		Outcome			GRA	ADE (outcome	e level)		
Interventions	Study	1	2	3	4	5	6	7		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade**
Multifactorial criteria-based algorithm vs.	Mendiguchia	?	×	×	?	✓	?	 ✓	Return to play	RCT ✓	?	?	√	xx	?	↓↓
lengthening hamstring exercises	et al. (5)	•							Reinjuries	RCT ✓	?	?	✓	xx	?	$\downarrow\downarrow$
Lengthening hamstring exercises versus to	Askling et al.	×	×	×	×	✓	?	?	Return to play		xx	✓	✓	✓	?	↓↓
conventional hamstring exercises (6)	(7,8)	×	×	×	×	✓	?	?	Reinjuries	RCT ✓	xx	✓	✓	××	?	$\downarrow\downarrow\downarrow$
Running and eccentric hamstring strengthening versus agility and trunk stabilization	Silder et al. (9)	?	?	×	?	✓	?	?	Return to play	RCT ✓	×	?	×	xx	?	\
Agility and trunk stabilization vs. hamstring	Sherry et al.	?	?	×	×	Ş	2	x	Return to play	RCT ✓	xx	?	×	×	?	$\downarrow\downarrow\downarrow\downarrow$
stretching and strengthening	(10)	·	ŗ	^					Reinjuries	RCT ✓	xx	?	×	×	?	$\downarrow\downarrow\downarrow$
Hamstring stretching four times/day versus hamstring stretching once daily	Malliaropoulos et al. (11)	?	?	×	?	?	?	×	Return to play	RCT ✓	xx	?	×	√	?	\
	Reurink et al. (12)	✓	✓	✓	✓	√	?	✓	Return to play	RCT ✓	✓	×	✓	✓	?	4
Platelet-rich plasma versus placebo or rehabilitation (6)	Hamilton et al. (13)	?	?	✓	✓	✓	×	✓		RC1 V	•	~	•	•	:	V
	Hamid et al. (14)	✓	✓	×	✓	✓	×	✓	Reinjuries	RCT ✓	✓	✓	✓	×	?	↓
Pain-threshold (≤4 on the 0-10 NRS) versus Pain-free (0 on the 0-10 NRS)	Hickey et al.	x	✓	/	/	2	2	/	Return to play	DOT 1	✓	?	×	×	?	↓↓
rehabilitation	(15)	^	_	_	•	:		_	Reinjuries	RCT √	√	?	√	XX	?	$\downarrow \downarrow$

Abbreviations: RCT (randomized controlled trial)

*Item 1: Random sequence generation; Item 2: Allocation concealment; Item 3: Blinding of participants and personal; Item 4: Blinding of outcome assessor; Item 5: Incomplete outcome data; Item 6: Selective reporting; Item 7: Other sources of bias.

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: x = item cause for possible downgrade once; xx = item cause for possible downgrade twice; x = item fulfilled, no downgrading; ? = item unclear or not available, no upgrading or downgrading.

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Table 5. Risk of bias asses	ssment and (GRAI	DE fo	or tre	atme	nt of	rectus	s fem	oris/c	uadr	iceps	injuri	es.										
Treatment Rectus femoris/	quadriceps						SI	GN C	heck	list 3'	k					Outcome			G	RADE (outcor	ne level)		
Interventions	Study	1	2	3	4	5	6	7	8	9	10	11	12	13	14		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade**
A two-phase criteria-based	Cross et al.	./	N/A	×	NI/A	2	N/A	×	×	~	2	\ \ \	N/A	2	~	Return to play	Cohort XX	×	?	×	×	?	1
intervention	(16)		IN/A	^	N/A		N/A	^	^	^	ŗ	^	N/A		_	Reinjuries	Cohort XX	×	?	✓	✓	?	1

Abbreviations: N/A (not applicable).

*Item 1: The study addresses an appropriate and clearly focused question?; Item 2: The two groups being studied are selected from source populations that are comparable in all respects other than the factor under investigation?; Item 3: The study indicates how many of the people asked to take part did so, in each of the groups being studied?; Item 4: The likelihood that some eligible subjects might have the outcome at the time of enrolment is assessed and taken into account in the analysis?; Item 5: What percentage of individuals or clusters recruited into each arm of the study dropped out before the study was completed?; Item 6: Comparison is made between full participants and those lost to follow up, by exposure status?; Item 7: The outcomes are clearly defined?; Item 8: The assessment of outcome is made blind to exposure status. If the study is retrospective this may not be applicable?; Item 9: Where blinding was not possible, there is some recognition that knowledge of exposure status could have influenced the assessment of outcome?; Item 10: The method of outcome assessment of exposure is reliable?; Item 11: Evidence from other sources is used to demonstrate that the method of outcome assessment is valid and reliable?; Item 12: Exposure level or prognostic factor is assessed more than once?; Item 13: The main potential confounders are identified and taken into account in the design and analysis?; Item 14: Have confidence intervals been provided?

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: x = item cause for possible downgrade once; xx = item cause for possible downgrade twice; x = item fulfilled, no downgrading; ?= item unclear or not available, no upgrading or downgrading.

** $\downarrow\downarrow\downarrow$ =downgrade quality by three levels

Table 6. Risk of bias asses	ssment and	GRA	DE fo	or tre	atme	nt of o	calf in	juries	3.														
Treatment Calf						;	SIGN	Che	cklist	3 and	d 4*					Outcome			G	RADE (outco	me level)		
Interventions	Study	1	2	3	4	5	6	7	8	9	10	11	12	13	14		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade***
M. daine and all two attendants are an area	Millow (4.7)	×	N//A	×	N/A	2	2	?	x		2		×	×	×	Return to play	Cohort ××	xx	?	✓	?	?	1
Multimodal treatment program	Millar (17)		N/A	•	N/A	!	!	!		N/A	:	N/A			•	Reinjuries	Cohort XX	xx	?	✓	?	?	1
Multimodal treatment program	Pedret et al. (18)	✓	N/A	x	N/A	✓	N/A	x	x	?	?	x	?	?	x	Reinjuries	Cohort XX	×	?	✓	?	?	1
Platelet-rich plasma**	Borrione et al. (19)	×	✓	✓	?	✓	✓	✓	×	?	x	×	-	-	-	Return to play	Case- control	xx	?	✓	✓	?	*

Abbreviations: N/A (not applicable).

*SIGN 3: Item 1: The study addresses an appropriate and clearly focused question?; Item 2: The two groups being studied are selected from source populations that are comparable in all respects other than the factor under investigation?; Item 3: The study indicates how many of the people asked to take part did so, in each of the groups being studied?; Item 4: The likelihood that some eligible subjects might have the outcome at the time of enrolment is assessed and taken into account in the analysis?; Item 5: What percentage of individuals or clusters recruited into each arm of the study dropped out before the study was completed?; Item 6: Comparison is made between full participants and those lost to follow up, by exposure status?; Item 7: The outcomes are clearly defined?; Item 8: The assessment of outcome is made blind to exposure status. If the study is retrospective this may not be applicable?; Item 9: Where blinding was not possible, there is some recognition that knowledge of exposure status could have influenced the assessment of outcome?; Item 10: The method of outcome assessment of exposure is reliable?; Item 11: Evidence from other sources is used to demonstrate that the method of outcome assessment is valid and reliable?; Item 12: Exposure level or prognostic factor is assessed more than once?; Item 13: The main potential confounders are identified and taken into account in the design and analysis?; Item 14: Have confidence intervals been provided?

SIGN 4: Item 1: The study addresses an appropriate and clearly focused question?; Item 2: The cases and controls are taken from comparable populations?; Item 3: The same exclusion criteria are used for both cases and controls?; Item 4: What percentage of each group (cases and controls) participated in the study?; Item 5: Comparison is made between participants and non-participants to establish their similarities or differences?; Item 6: Cases are clearly defined and differentiated from controls?; Item 7: It is clearly established that controls are non-cases?; Item 8: Measures will have been taken to prevent knowledge of primary exposure influencing case ascertainment?; Item 9: Exposure status is measured in a standard, valid and reliable way?; Item 10: The main potential confounders are identified and taken into account in the design and analysis?; Item 11: Confidence intervals are provided.

** Risk of bias using SIGN 4

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: × = item cause for possible downgrade once; ×× = item cause for possible downgrade twice; ✓ = item fulfilled, no downgrading; ?= item unclear or not available, no upgrading or downgrading.

*** $\downarrow \downarrow \downarrow$ =downgrade quality by three levels

Supplementary material

Table 7. Risk of bias assessment and Prevention Hamstring							ent Item	ו*	Outcome			GR <i>A</i>	DE (outcome le	evel)		
Interventions	Study	1	2	3	4	5	6	7		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade
	Gabbe et al. (21)	✓	✓	x	x	✓	✓	✓								
	Soligard et al. (22)	?	?	×	×	×	✓	✓								
	Engebretsen et al. (23)	?	?	×	×	?	?	×								
Interventions including the Nordic Hamstring exercise (20)	Petersen et al. (24)	✓	√	×	×	✓	✓	✓	Injuries	RCT ✓	?/√	×	✓	✓	✓	\
	Del Ama Espinosa et al. (25)	✓	✓	×	×	✓	✓	✓								
	Silvers-Granelli et al. (26)	✓	?	×	x	×	?	x								
	Van der Horst et al. (27)	✓	✓	×	×	✓	✓	✓								
	Aksling et al. (29)	?	?	x	?	?	?	×								
Mixed eccentric hamstring training (28)	Gabbe et al. (21)	✓	✓	×	x	✓	✓	✓	Injuries	RCT✓	✓	×	✓	×	?	↓ ↓
	Engebretsen et al. (23)	?	?	×	×	?	?	×	_						•	
	Petersen et al. (24)	✓	✓	×	×	✓	✓	✓								
FIFA 11+ (30)	Soligard et al. (22)	?	?	×	×	×	✓	✓	. Injuries	RCT ✓	x	✓	✓	✓	?	\
, ,	Silvers-Granelli et al. (26)	✓	?	×	×	×	?	×	_	1.01					•	
Nordic Hamstring Exercise Protocol (meta- analysis performed as part of this	Petersen et al. (24)	✓	✓	×	×	✓	✓	✓	Injuries	DOT V	√	✓	✓	√	?	\leftrightarrow
statement)	Van der Horst et al. (27)	✓	✓	×	×	✓	✓	✓	iiijulies	RCT ✓	•	•	•	•		↔
Bounding exercise program	Van de Hoef et al. (31)	✓	?	×	×	✓	✓	✓	Injuries	RCT ✓	✓	?	✓	×	?	\
FIFA 11+ program pre- and post-football	Al Attar et al. (32)	✓	?	x	×	✓	✓	×	Injuries	RCT ✓	x	?	✓	xx	?	1
Modified FIFA 11+ with rescheduling of Part 2 versus standard FIFA 11+	Whalan et al. (33)	?	?	x	×	✓	?	✓	Injuries	RCT ✓	?	?	✓	×	?	\
Balance board training	Soderman et al. (34)	?	?	×	×	✓	?	×	Injuries	RCT ✓	×	?	✓	xx	?	$\downarrow\downarrow\downarrow$

Abbreviations: RCT (randomized controlled trial)

*Item 1: Random sequence generation; Item 2: Allocation concealment; Item 3: Blinding of participants and personal; Item 4: Blinding of outcome assessor; Item 5: Incomplete outcome data; Item 6: Selective reporting; Item 7: Other sources of bias.

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

Supplementary material

GRADE assessments: x = item cause for possible downgrade once; x = item cause for possible downgrade twice; x = item fulfilled, no downgrading; ? = item unclear or not available, no upgrading or downgrading.

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Table 8. Risk of bias assessment an		reventi		<u> </u>	•											
Prevention Adductor (Gro							nt Item*		Outcome				ADE (outcome le			
Interventions	Study	1	2	3	4	5	6	7		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade **
	Arnason et al. (36)	?	?	×	×	?	?	×								
	Beijsterveldt et al. (37)	?	?	×	x	✓	×	?								
Mixed groin prevention programs (35)	Engebretsen et al. (23)	?	?	x	×	?	?	×	Injuries	RCT √	×	✓	×	×	?	*
winced grown prevention programs (65)	Holmich et al. (38)	✓	✓	×	×	✓	?	×	injunes	HC1					•	•
	Soderman et al. (34)	?	?	×	×	✓	?	×								
	Steffen et al. (39)	?	?	×	×	✓	?	×								
Specific adductor strength training (35)	Holmich et al. (38)	✓	✓	×	x	✓	?	×	Injuries	DOT 1	×	✓	×	x	?	*
Specific adductor strength training (55)	Engebretsen et al. (23)	?	?	x	×	?	?	×	irijuries	RCT ✓	^	•	^	~		$\Psi\Psi$
EIEA 44 (OE)	Steffen et al. (39)	?	?	x	×	✓	?	×			x	×	×	×	?	
FIFA 11 (35)	Beijsterveldt et al. (37)	?	?	×	×	✓	×	?	Injuries	RCT ✓		*		*	:	$\downarrow\downarrow\downarrow$
FIFA 11+ programme in footabll (30)	Silvers- Granelli et al. (26)	✓	?	x	×	x	?	×	Injuries	RCT √	×	✓	×	√	ç	↓ ↓
	Soligard et al. (22)	?	?	×	×	×	✓	✓	-							
FIFA 11+ programme in mixed sports	Longo et al. (40)	\checkmark	?	×	x	✓	✓	✓	Injuries	RCT ✓	✓	×	×	xx	· P	$\downarrow\downarrow\downarrow\downarrow$
THE Programme in mixed sports	Slauterbeck et al. (41)	✓	?	×	×	✓	?	✓	injunes	HUI *	•				•	***
Adductor strengthening program	Haroy et al. (42)	✓	?	×	×	✓	✓	✓	Injuries	RCT ✓	✓	?	×	✓	?	\
FIFA 11+ program pre- and post-football	Al Attar et al. (32)	✓	?	×	×	✓	✓	×	Injuries	RCT ✓	×	?	×	xx	?	1
Modified FIFA 11+ with rescheduling of Part 2 versus standard FIFA 11+	Whalan et al. (33)	?	?	×	×	✓	?	✓	Injuries	RCT ✓	?	?	×	✓	?	\

Abbreviations: RCT (randomized controlled trial)

*Item 1: Random sequence generation; Item 2: Allocation concealment; Item 3: Blinding of participants and personal; Item 4: Blinding of outcome assessor; Item 5: Incomplete outcome data; Item 6: Selective reporting; Item 7: Other sources of bias.

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: \times = item cause for possible downgrade once; \times = item cause for possible downgrade twice; \checkmark = item fulfilled, no downgrading; ? = item unclear or not available, no upgrading or downgrading.

** \downarrow = downgrade quality by one level; \downarrow \downarrow =downgrade quality by two levels; \downarrow \downarrow \downarrow =downgrade quality by three levels; \leftrightarrow =no downgrade

Table 9. Risk of bias assessment and	d GRADE for p	reventi	on of	anterio	r thigh	n/quad	driceps i	njuries.								
Prevention anterior thigh/quad	driceps		Risk	of Bia	s asse	essme	nt Item		Outcome			GRA	ADE (outcome l	evel)		
Interventions	Study	1	2	3	4	5	6	7		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade
FIFA 11+ (meta-analysis performed as part	Silvers- Granelli et al. (26)	✓	?	×	x	x	?	×	Injuries	RCT √	×	✓	✓	×	?	4
of this statement)	Soligard et al. (22)	?	?	x	x	x	✓	✓	, ,	nor -					•	,
FIFA 11+ program pre- and post-football	Al Attar et al. (32)	✓	?	×	×	✓	✓	×	Injuries	RCT ✓	×	?	✓	××	?	$\downarrow\downarrow\downarrow\downarrow$
Modified FIFA 11+ with rescheduling of Part 2 versus standard FIFA 11+	Whalan et al. (33)	?	?	×	×	✓	?	✓	Injuries	RCT ✓	?	?	✓	✓	?	\leftrightarrow
Balance board training	Soderman et al. (34)	?	?	×	×	✓	?	×	Injuries	RCT ✓	×	?	✓	××	?	1

Abbreviations: RCT (randomized controlled trial)

*Item 1: Random sequence generation; Item 2: Allocation concealment; Item 3: Blinding of participants and personal; Item 4: Blinding of outcome assessor; Item 5: Incomplete outcome data; Item 6: Selective reporting; Item 7: Other sources of bias.

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: \times = item cause for possible downgrade once; \times = item cause for possible downgrade twice; \checkmark = item fulfilled, no downgrading; ? = item unclear or not available, no upgrading or downgrading.

^{**} \downarrow = downgrade quality by one level; \downarrow \downarrow =downgrade quality by two levels; \downarrow \downarrow \downarrow =downgrade quality by three levels; \leftrightarrow =no downgrade

Table 10. Risk of bias asse	essment and (GR/	ADE	for p	orever	ntion	of cal	f inju	ries.														
Prevention Ca	lf						S	SIGN	Chec	klist 3	}*					Outcome			G	RADE (outco	me level)		
Interventions	Study	1	2	3	4	5	6	7	8	9	10	11	12	13	14		Study design	Risk of bias	Inconsistency	Indirectness	Imprecise evidence	Publication bias	Downgrade**
soccer-specific balance program	Kraemer et al. (43)	✓	N/ A	N/A	?	×	?	✓	x	x	✓	✓	?	?	×	Injuries	Cohort XX	×	?	✓	?	?	\

Abbreviations: N/A (not applicable).

*SIGN 3: Item 1: The study addresses an appropriate and clearly focused question?; Item 2: The two groups being studied are selected from source populations that are comparable in all respects other than the factor under investigation?; Item 3: The study indicates how many of the people asked to take part did so, in each of the groups being studied?; Item 4: The likelihood that some eligible subjects might have the outcome at the time of enrolment is assessed and taken into account in the analysis?; Item 5: What percentage of individuals or clusters recruited into each arm of the study dropped out before the study was completed?; Item 6: Comparison is made between full participants and those lost to follow up, by exposure status?; Item 7: The outcomes are clearly defined?; Item 8: The assessment of outcome is made blind to exposure status. If the study is retrospective this may not be applicable?; Item 9: Where blinding was not possible, there is some recognition that knowledge of exposure status could have influenced the assessment of outcome?; Item 10: The method of outcome assessment of exposure is reliable?; Item 11: Evidence from other sources is used to demonstrate that the method of outcome assessment is valid and reliable?; Item 12: Exposure level or prognostic factor is assessed more than once?; Item 13: The main potential confounders are identified and taken into account in the design and analysis?; Item 14: Have confidence intervals been provided?

Risk of bias assessment: × item not fulfilled; ✓ = item fulfilled; ? unclear or unknown if item is fulfilled

GRADE assessments: x = item cause for possible downgrade once; xx = item cause for possible downgrade twice; x = item fulfilled, no downgrading; ?= item unclear or not available, no upgrading or downgrading.

** $\downarrow\downarrow\downarrow$ =downgrade quality by three levels

ROBIS: Tool to assess risk of bias in systematic reviews

Table 11. Suggested Tabular Presentation for ROBIS Results

Review		Phase 2			Phase 3
_	1. STUDY ELIGIBILITY CRITERIA	2. IDENTIFICATION AND SELECTION OF STUDIES	3. DATA COLLECTION AND STUDY APPRAISAL	4. SYNTHESIS AND FINDINGS	RISK OF BIAS IN THE REVIEW
Van Dyk 2019 (20)	<u>©</u>	8	<u>©</u>	?	?
Thorborg 2017 (30)			\odot	?	?
Esteve 2015 (35)	?	?	<u>©</u>	?	?
Goode 2015 (28)	?		?	?	?
Pas 2015 (6)	?	<u>©</u>	<u>©</u>	?	?
Rieman 2013 (44)	?	?	©	?	?

© = low risk; <mark>⊖ =</mark> high risk; ? = unclear risk

Supplementary material

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